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INTELLECTUAL PROPERTY LAW
(PATENT, BIOTECHNOLOGY, COMPUTER,
TRADEMARK & TRADE SECRET LAW)

jc558 U.S. PTO
09/296031
04/21/99

April 21, 1999

Docket No.: D6218

The Assistant Commissioner Of Patents And Trademarks
BOX PATENT APPLICATION
Washington, DC 20231

Dear Sir:

Transmitted herewith for filing is the non-provisional patent application which is attached hereto in the:

Name of: **Susan A. Lyons and Harald W. Sontheimer**
For: *Diagnosis and Treatment of Neuroectodermal Tumors*

CLAIMS AS FILED

<u>Fee for:</u>	<u>Small entity</u>	<u>Amount</u>
Basic fee	\$ 3 8 0	\$ 3 8 0
Each independent claim in excess of 3 (1)		
Each claim excess of 20 (0)		
Multiple dependent claim		

TOTAL FILING FEE \$ 3 8 0

___ Please charge my Deposit Account No. _____ in the total amount of the filing fee and the assignment recordation fee if any.

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X Any additional fees under 37 CFR 1.16.
X Any application processing fees under 37 CFR 1.17.

X Small Entity Statement

A small entity statement is enclosed and its benefit under 37 CFR 1.28(a) is hereby claimed.

Relate Back--35 U.S.C. 119(e)

This non-provisional application claims benefit of priority of provisional application filed

Assignment

The application is assigned by the inventors to

Sequence Listing

The sequence listing is enclosed, including a paper copy, a copy on disk for the computer readable form and a compliance letter indicating that the sequence listing on the paper copy and the disk are one and the same.

X Power of Attorney

X is attached.

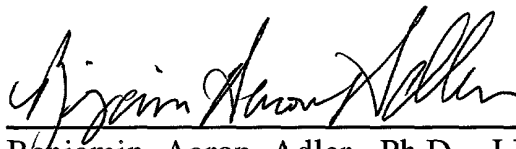
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X Two photocopies of this sheet are enclosed.

Date:

April 21, 1999



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Applicant or Patentee: Susan Lyons + Harald W. Sonthheimer Attorney's
Serial or Patent No.: _____ Docket No.: 6218
Filed or Issued: April 21, 1999
For: Diagnosis and Treatment of Neuroectodermal Tumors

VERIFIED STATEMENT (DECLARATION) CLAIMING SMALL ENTITY
STATUS (37 CFR 1.9(f) and 1.27(c)) - NONPROFIT ORGANIZATION

I hereby declare that I am an official of the nonprofit organization empowered to act on behalf of the concern identified below:

NAME OF ORGANIZATION University of Alabama at Birmingham Research Foundation
ADDRESS OF CONCERN 701 20th Street South, Birmingham, AL 35294-0011
X University or other institution of higher education

I hereby declare that the above identified nonprofit organization qualifies as a nonprofit organization as defined in 37 CFR 1.9(e), for purposes of paying reduced fees under section 41(a) and (b) of Title 35, United States Code.

I hereby declare that rights under contract or law have been conveyed to and remain with the nonprofit organization identified above with regard to the invention, entitled as above

_____ by inventor(s) Susan Lyons
and Harald W. Sonthheimer described in:
☒ the specification filed herewith
☐ application serial no. _____, filed
☐ patent no. _____, issued

If the rights held by the above identified nonprofit organization are not exclusive, each individual, concern or organization having rights to the invention is listed below* and no rights to the invention are held by any person, other than the inventor, who could not qualify as a small business concern under 37 CFR 1.9(d) or by any concern which would not qualify as a small business concern under 37 CFR 1.9(d) or a nonprofit organization under 37 CFR 1.9(e).

*NOTE: Separate verified statements are required from each named person, concern or organization having rights to the invention averring to their status as small entities. (37 CFR 1.27)

NAME

ADDRESS

☐ INDIVIDUAL

☐ SMALL BUSINESS CONCERN

☐ NONPROFIT ORGANIZATION

I acknowledge the duty to file, in this application or patent, notification of any change in status resulting in loss of entitlement to small entity status prior to paying, or at the time of paying, the earliest of the issue fee or any maintenance fee due after the date on which status as a small entity is no longer appropriate. (37 CFR 1.28(b))

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon, or any patent to which this verified statement is directed.

NAME OF PERSON SIGNING David L. Day
TITLE OF PERSON OTHER THAN OWNER Interim Director, UAB Research Foundation
SIGNATURE David L. Day
DATE 4/21/99

DIAGNOSIS AND TREATMENT OF NEUROECTODERMAL TUMORS

5

10

BACKGROUND OF THE INVENTION

Federal Funding Legend

15

This invention was produced in part using funds from the Federal government under NIH grant no. R01 NS 36692. Accordingly, the Federal government has certain rights in this invention.

Field of the Invention

20

The present invention relates generally to the fields of cell physiology, neurology, developmental biology, and oncology. More specifically, the present invention relates to novel methods of

using a chlorotoxin sensitive cytoplasmic protein for the diagnosis and treatment of primitive neuroectodermal tumors (PNET).

Description of the Related Art

5 During embryonic development, the future nervous system forms from a specialized layer of ectodermal cells called the neuroectoderm. This layer extends longitudinally along the body axis congruent with the future spinal column. Invagination of the neuroectoderm gives rise to the neural tube from which essentially
10 all central nervous system (CNS) components including the spinal cord develop. Specialized cell clusters along the rim of the invaginating neural tube stay separate from the tube and from the neural crest. These highly migratory neuroectodermal cells give rise to specialized cells throughout the body including Schwann cells,
15 neuronal cells of the peripheral nervous system (PNS) (enteric, parasympathetic, sympathoadrenal, and sensory neurons), pigment cells (melanocytes), endocrine cells and cells forming connective tissue of the face and neck. Since these cells share a common embryonic origin with central nervous system cells, it is not
20 surprising that these cells, or the tumors developing from these cells,

share some genetic and antigenic phenotypes with central nervous system cells.

For example, melanomas and glioblastomas share a common mutation in the gene encoding for the epidermal growth factor receptor (EGFR) (1). Malignant astrocytomas and neurofibromas not only express high levels of the epidermal growth factor receptors but also vascular endothelial growth factor receptor (VEGF-R) and platelet-derived growth factor receptor (PDGF-R) (2).

A high expression of the mutant variant, EGFRvIII, has been demonstrated in glial tumors as well as the extracellular matrix proteins, GP 240 and tenascin (3). Tenascin and the ganglioside-3',6'-isoLD1 have been found in both gliomas and primitive neuroectodermal tumor tissues (3). In another study, antibodies to tenascin bound extensively to CNS gliomas and also to melanomas, breast, lung and squamous cell carcinomas (4). Another ganglioside, GD2, has been shown to be a common antigen marker in both gliomas and primitive neuroectodermal tumor tissues (5). Other common antigens between melanomas and gliomas were demonstrated by showing that Tyr, TRP-1, TRP-2 and gp100 gene products are commonly found in both melanoma and glioma tumors (6). Common cytokines or their receptors linking tumors of astrocytomas,

ependymomas and primitive neuroectodermal tumors have been identified as: interleukin (IL) IL-1 alpha, IL-1, IL-1R1, IL-1R antagonist and transforming growth factor (TGF) TGF-beta 1 (11).

Another class of proteins used as markers for gliomas and primitive neuroectodermal tumors are the cytoskeletal proteins, neurofilament (NF), glial fibrillary acidic protein (GFAP), intermediate filaments (IF), intermediate associated protein filament (IFAP), vimentin, nestin and keratins. These markers have been used to determine stages of differentiation along the various cell lineages (12). New evidence linking astrocytomas with certain primitive neuroectodermal tumor tumors is the cytoskeleton marker of IFAP-300kDa, a marker of immature glia (13).

Further arguments for a tight linkage of neuroectodermally derived cells in the central nervous system and periphery can be made based on their similar dependence on epigenetic influences. For example, sympathoadrenal precursor neurons require basic fibroblast growth factor (bFGF) to proliferate and differentiate, but survival of these cells depends on nerve growth factor (NGF) responsiveness and nerve growth factor availability (14). A similar scenario is required for each of the other cell types. Not only are growth and trophic factors necessary but

cytokines and hormones are needed for which links remain to be elucidated between primitive neuroectodermal tumors and gliomas.

However, despite this list of similarities shared between neuroectodermally derived cells, these cells are distinct entities with unique cytological, biochemical and functional features. Indeed, the list of unique features not shared with other neuroectodermally derived cells by far exceeds the above mentioned shared phenotypes. Thus, one can not assume *a priori* that expression of a certain antigen or phenotype is to be expected in a given cell type based on expression by any other member of the neuroectodermally derived cell types.

Neuroblastomas generally express a selective increase in the gene copy number of the MYCN gene found in fetal stages of brain development suggesting links between the origin of the cells and the ability of neoplastic cells to dedifferentiate (7). However, this gene has yet to be demonstrated in the glioma cells. Other proteins that are not common to both glioma and primitive neuroectodermal tumors have been demonstrated. CD99 immunoreactivity is used as a tool in identifying primitive neuroectodermal tumors (8) and has been shown in Ewing's sarcoma tumors although not in gliomas (9). Another factor, stem cell factor

and its receptor, c-kit, are also expressed in both primitive neuroectodermal tumor and Ewing's Sarcoma tumors (10).

5 The common origin and ability to respond to internal and external signals during the normal developmental processes suggests that central nervous system cells and peripheral neuroectodermally derived cells may also share common mechanisms during pathological developments as for example, during neoplasia. Such neoplastic tissues include CNS gliomas that are glial-derived tumor cells specific to the CNS. They metastasize only within the CNS
10 including the spinal column. They are believed to originate from at least three separate lineages either from undifferentiated precursor cells or by dedifferentiation of astrocytes, oligodendrocytes or ependymal cells.

15 Primitive neuroectodermal tumors (PNET) are found both in the CNS and PNS. Primitive neuroectodermal tumors found only in the PNS are referred to as peripheral primitive neuroectodermal tumors (PPNET). Primitive neuroectodermal tumors manifest preferentially in children and have capacity for developing into a variety of neuronal, astrocytic, ependymal, muscular and melanotic
20 lines. The conceptual basis of grouping these tumors together is based upon sharing common progenitor cells as well as sharing

similar neoplastic transformations leading to tumors of similar morphological features and biological behavior. However, there remains controversy in placing all primitive neuroectodermal tumors into the same categories. The following paragraphs demonstrate
5 examples of the overlap of common antigens between the various types of CNS and PNS tumors.

Supratentorial primitive neuroectodermal tumors include cerebral medulloblastomas, cerebral neuroblastomas, 'blue' tumors, ependymoblastoma and other primitive neuroectodermal tumors,
10 such as pineoblastomas (WHO grade IV). The most useful markers for these tumors include GFAP, NFP, desmin and melanin. Others antigens found in these tumors are vimentin, nestin, keratin but are not useful for diagnostic purposes.

Peripheral neuroblastic tumors of the adrenal gland
15 (medulla) and sympathetic nervous system are the most common type of childhood tumor outside of the CNS. Primary sites for these primitive neuroectodermal tumors are in the adrenals, abdominal, thoracic, cervical and pelvic sympathetic ganglia but include other primary sites as orbit, kidney, lung, skin, ovary, spermatic cord, and
20 urinary bladder. Specific names of these related tumors are pheochromocytomas, paraganglioma, neuroblastomas,

ganglioneuromas, ganglioneuroblastomas, neurofibromas, schwannomas, and malignant peripheral nerve sheath tumors. These all share common origin in the neural crest. Neuroblastomas all share high TRK-A (NGFR) and CD44 expressions. Neuronal specific enolase (NSE), synaptophysin, neural filament (NF) protein, GD2, tyrosine hydroxylase (TH) and chromogranin are used as diagnostic markers also found in medulloblastomas. Neuroblastomas generally express a selective increase in the gene copy number of the *MYCN* gene found in fetal stages of brain development (7).

Medulloblastomas are members of the primitive neuroectodermal tumors that are described as highly malignant embryonal tumors of the CNS found in the cerebellum (WHO grade IV). A common antigen of these medulloblastoma and other neuronal lineage tumors is synaptophysin (not found in glial or mesenchymal brain tumors). Nestin (IF protein) is found in developing CNS precursor cells and in medulloblastomas and in some peripheral neuroectodermal origin cells. Nestin (and vimentin) are found in medulloblastomas, astrocytomas, glioblastomas, ependymomas, gangliogliomas and meningiomas (only GFAP is found in the astrocytic-derived cells, which are occasionally 'trapped' in medulloblastomas). Increased levels of neural-cellular adhesion

molecule (N-CAM) found in these tumors, may reflect levels of differentiation in the development of tumors (15). While varying levels of nerve growth factor (NGF), are found in nearly all tumors, medulloblastomas exhibited substantial reactivity to the NGF receptor and related proteins, neurotrophin (NT) NT-3, TRK-C and brain derived neurotrophic factor (BDNF) (16).

Melanomas, arising from melanocytes follow a graded development from diffuse melanocytosis, to melanocytoma to malignant melanomas. S100 protein is a marker for these tumors, as vimentin and NSE reactivity are variable.

Small cell neuroendocrine carcinomas of the lung are highly invasive and typically found in adult smokers. They have been shown to exhibit reactivity to many of the neural and neuroendocrine markers (some of them similar to N-CAMs) for tumor differentiation as peripheral primitive neuroectodermal tumors, gliomas, and ependymomas. These markers include neural specific enolase and extremely high *c-src* expression (17).

A feature conspicuously shared between developing CNS cells and neural crest derived cells is their propensity to migrate either towards a target or target area. It is believed that this ability is lost after cell differentiation and maturation. However, tumors of

the CNS show significant cell migration and invasion into healthy brain, suggesting that cell have maintained or regained this enhanced migratory ability. It is, thus, not surprising that neoplastic transformation of neuroectodermally derived cells outside the CNS would have similar migratory abilities. At the intended destination, these cells differentiate into their final phenotype, similar to normal development, influenced by several trophic factors crucial for the proliferation and differentiation of various cell types.

The prior art is deficient in the lack of an diagnostic and therapeutic agents specifically targeted to primitive neuroectodermal tumors. The present invention fulfills this longstanding need and desire in the art.

SUMMARY OF THE INVENTION

In one embodiment of the current invention, a method is described for the treatment of tumors of neuroectodermal origin by administering a ligand specific for this class of tumors fused to a cytotoxic moiety. Specific neuroectodermal tumor tumors which can

be treated in this manner include gliomas, meningiomas, ependymomas, medulloblastomas, neuroblastomas, gangliomas, pheochromocytomas, melanomas, peripheral primitive neuroectodermal tumors, small cell carcinoma of the lung, Ewing's sarcoma, and metastatic tumors of neuroectodermal origin in the brain.

In the preferred embodiment, the neuroectodermal tumor specific ligand is chlorotoxin in the form of a fusion protein of chlorotoxin with cytotoxic moiety. The chlorotoxin may be native, synthetic or recombinant chlorotoxin. Possible cytotoxic moieties include gelonin, ricin, saponin, pseudomonas exotoxin, pokeweed antiviral protein, diphtheria toxin, and complement proteins.

In another embodiment of the current invention, the neuroectodermal tumor specific ligand is an antibody against the chlorotoxin receptor, presumably a 72 kDa chloride channel. The antibody may be fused to gelonin, ricin, saponin, pseudomonas exotoxin, pokeweed antiviral protein, diphtheria toxin, and complement proteins.

In yet another embodiment of the current invention, a method of differentiating neuroectodermal tumor-derived neoplastic tumor tissue from non-neoplastic tissue is presented. This is

accomplished by exposing the tissue with labeled chlorotoxin and measuring the binding of the labeled chlorotoxin. An elevated level of binding relative to normal tissue is indicative that the tissue is neoplastic. In one embodiment, the label is a fluorescent moiety which is detected by fluorescent microscopy, fluorescent activated cell sorting or a fluorescent plate reader. Alternatively, the chlorotoxin may be radiolabeled (eg. ^{131}I -chlorotoxin or ^{125}I -chlorotoxin; a person having ordinary skill in this art would readily recognize other useful radiolabels) and detected by positron emission tomography scanning. Alternatively, the chlorotoxin may be conjugated to a non-fluorescent detection moiety such as biotin and detected immunohistochemically or by use of a colorimetric assay.

BRIEF DESCRIPTION OF THE DRAWINGS

So that the matter in which the above-recited features, advantages and objects of the invention, as well as others which will become clear, are attained and can be understood in detail, more particular descriptions of the invention briefly summarized above may be had by reference to certain embodiments thereof which are

illustrated in the appended drawings. These drawings form a part of the specification. It is to be noted, however, that the appended drawings illustrate preferred embodiments of the invention and therefore are not to be considered limiting in their scope.

5 **Figure 1** shows the positive immunohistochemical staining of a glioblastoma multiform (GBM) tumor with chlorotoxin. The brown reaction product of DAB 3'3'-diaminobenzidine with biotinylated chlorotoxin is clearly visible in the TM-601 stained section. TM-601: biotinylated chlorotoxin stained, counter-stained
10 with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

Figure 2 demonstrates that normal brain is not immunohistochemically stained by biotinylated chlorotoxin. TM-601: biotinylated chlorotoxin, counter-stained with methyl green; Control:
15 methyl green only; and, H&E stain: hematoxylin and eosin. Normal brain was also stained with biotinylated antibodies against GFAP (Glial Fibrillary Acidic Protein) which positively stains astrocytes in normal brain tissue.

Figure 3 shows chlorotoxin staining of an adrenal mass
20 neuroblastoma tumor. TM-601: biotinylated chlorotoxin, counter-

stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

Figure 4 illustrates that biotinylated chlorotoxin immunohistochemically stains pheochromocytomas. TM-601:

5 biotinylated chlorotoxin, counter-stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

Figure 5 illustrates that normal adrenal tissue is not immunohistochemically stained by biotinylated chlorotoxin. TM-601:

10 biotinylated chlorotoxin, counter-stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

Figure 6 shows the immunohistochemical staining with biotinylated chlorotoxin of melanoma tumor cells metastasized to the brain TM-601: biotinylated chlorotoxin, counter-stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

Figure 7 illustrates biotinylated chlorotoxin immunohistochemical staining of melanoma tumor cells metastasized to the lung. TM-601: biotinylated chlorotoxin, counter-stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

Figure 8 shows immunohistochemical staining of normal skin with biotinylated chlorotoxin. TM-601: biotinylated chlorotoxin, counter-stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

5 **Figure 9** shows immunohistochemical staining of small cell lung carcinomas with biotinylated chlorotoxin. TM-601: biotinylated chlorotoxin, counter-stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

10 **Figure 10** shows that biotinylated chlorotoxin does not immunohistochemically stain normal lung tissue. TM-601: biotinylated chlorotoxin, counter-stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

15 **Figure 11** shows the immunohistochemical staining of a medulloblastoma tumor with biotinylated chlorotoxin. TM-601: biotinylated chlorotoxin, counter-stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

20 **Figure 12** shows that the rare, neuroectodermally derived bone cancer, Ewing's sarcoma, also exhibits positive immunohistochemical staining with biotinylated chlorotoxin. TM-601: biotinylated chlorotoxin, counter-stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

5 **Figure 13** shows the negative results obtained in the immunohistochemical staining of normal stomach tissue with biotinylated chlorotoxin. TM-601: biotinylated chlorotoxin, counter-stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

Figure 14 shows the lack of immunohistochemical staining of normal liver tissue with biotinylated chlorotoxin. TM-601: biotinylated chlorotoxin, counter-stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

10 **Figure 15** demonstrates that normal spleen tissue is not immunohistochemically stained by with biotinylated chlorotoxin. TM-601: biotinylated chlorotoxin, counter-stained with methyl green; Control: methyl green only; and, H&E stain: hematoxylin and eosin.

15 DETAILED DESCRIPTION OF THE INVENTION

In accordance with the present invention there may be employed conventional molecular biology, microbiology, and recombinant DNA techniques within the skill of the art. Such

techniques are explained fully in the literature. See, e.g., Maniatis, Fritsch & Sambrook, "Molecular Cloning: A Laboratory Manual (1982); "DNA Cloning: A Practical Approach," Volumes I and II (D.N. Glover ed. 1985); "Oligonucleotide Synthesis" (M.J. Gait ed. 1984);
5 "Nucleic Acid Hybridization" [B.D. Hames & S.J. Higgins eds. (1985)]; "Transcription and Translation" [B.D. Hames & S.J. Higgins eds. (1984)]; "Animal Cell Culture" [R.I. Freshney, ed. (1986)]; "Immobilized Cells And Enzymes" [IRL Press, (1986)]; B. Perbal, "A Practical Guide To Molecular Cloning" (1984).

10 Therefore, if appearing herein, the following terms shall have the definitions set out below.

As used herein, the term "cDNA" shall refer to the DNA copy of the mRNA transcript of a gene.

As used herein, the term "derived amino acid sequence"
15 shall mean the amino acid sequence determined by reading the triplet sequence of nucleotide bases in the cDNA.

As used herein the term "screening a library" shall refer to the process of using a labeled probe to check whether, under the appropriate conditions, there is a sequence complementary to the
20 probe present in a particular DNA library. In addition, "screening a library" could be performed by PCR.

As used herein, the term "PCR" refers to the polymerase chain reaction that is the subject of U.S. Patent Nos. 4,683,195 and 4,683,202 to Mullis, as well as other improvements now known in the art.

5 The amino acid described herein are preferred to be in the "L" isomeric form. However, residues in the "D" isomeric form can be substituted for any L-amino acid residue, as long as the desired functional property of immunoglobulin-binding is retained by the polypeptide. NH₂ refers to the free amino group present at
10 the amino terminus of a polypeptide. COOH refers to the free carboxy group present at the carboxy terminus of a polypeptide. In keeping with standard polypeptide nomenclature, *J Biol. Chem.*, 243:3552-59 (1969), abbreviations for amino acid residues are known in the art.

15 It should be noted that all amino-acid residue sequences are represented herein by formulae whose left and right orientation is in the conventional direction of amino-terminus to carboxy-terminus. Furthermore, it should be noted that a dash at the beginning or end of an amino acid residue sequence indicates a
20 peptide bond to a further sequence of one or more amino-acid residues.

A "replicon" is any genetic element (e.g., plasmid, chromosome, virus) that functions as an autonomous unit of DNA replication *in vivo*; i.e., capable of replication under its own control.

A "vector" is a replicon, such as plasmid, phage or cosmid,
5 to which another DNA segment may be attached so as to bring about the replication of the attached segment.

A "DNA molecule" refers to the polymeric form of deoxyribonucleotides (adenine, guanine, thymine, or cytosine) in its either single stranded form, or a double-stranded helix. This term
10 refers only to the primary and secondary structure of the molecule, and does not limit it to any particular tertiary forms. Thus, this term includes double-stranded DNA found, *inter alia*, in linear DNA molecules (e.g., restriction fragments), viruses, plasmids, and chromosomes. In discussing the structure herein according to the
15 normal convention of giving only the sequence in the 5' to 3' direction along the nontranscribed strand of DNA (i.e., the strand having a sequence homologous to the mRNA).

An "origin of replication" refers to those DNA sequences that participate in DNA synthesis.

20 A DNA "coding sequence" is a double-stranded DNA sequence which is transcribed and translated into a polypeptide *in*

vivo when placed under the control of appropriate regulatory sequences. The boundaries of the coding sequence are determined by a start codon at the 5' (amino) terminus and a translation stop codon at the 3' (carboxyl) terminus. A coding sequence can include, but is not limited to, prokaryotic sequences, cDNA from eukaryotic mRNA, genomic DNA sequences from eukaryotic (e.g., mammalian) DNA, and even synthetic DNA sequences. A polyadenylation signal and transcription termination sequence will usually be located 3' to the coding sequence.

Transcriptional and translational control sequences are DNA regulatory sequences, such as promoters, enhancers, polyadenylation signals, terminators, and the like, that provide for the expression of a coding sequence in a host cell.

A "promoter sequence" is a DNA regulatory region capable of binding RNA polymerase in a cell and initiating transcription of a downstream (3' direction) coding sequence. For purposes of defining the present invention, the promoter sequence is bound at its 3' terminus by the transcription initiation site and extends upstream (5' direction) to include the minimum number of bases or elements necessary to initiate transcription at levels detectable above background. Within the promoter sequence will be

found a transcription initiation site, as well as protein binding domains (consensus sequences) responsible for the binding of RNA polymerase. Eukaryotic promoters often, but not always, contain "TATA" boxes and "CAT" boxes. Prokaryotic promoters contain Shine-Dalgarno sequences in addition to the -10 and -35 consensus sequences.

An "expression control sequence" is a DNA sequence that controls and regulates the transcription and translation of another DNA sequence. A coding sequence is "under the control" of transcriptional and translational control sequences in a cell when RNA polymerase transcribes the coding sequence into mRNA, which is then translated into the protein encoded by the coding sequence.

A "signal sequence" can be included near the coding sequence. This sequence encodes a signal peptide, N-terminal to the polypeptide, that communicates to the host cell to direct the polypeptide to the cell surface or secrete the polypeptide into the media, and this signal peptide is clipped off by the host cell before the protein leaves the cell. Signal sequences can be found associated with a variety of proteins native to prokaryotes and eukaryotes.

The term "oligonucleotide", as used herein in referring to the probe of the present invention, is defined as a molecule

comprised of two or more ribonucleotides, preferably more than three. Its exact size will depend upon many factors which, in turn, depend upon the ultimate function and use of the oligonucleotide.

The term "primer" as used herein refers to an oligonucleotide, whether occurring naturally as in a purified restriction digest or produced synthetically, which is capable of acting as a point of initiation of synthesis when placed under conditions in which synthesis of a primer extension product, which is complementary to a nucleic acid strand, is induced, i.e., in the presence of nucleotides and an inducing agent such as a DNA polymerase and at a suitable temperature and pH. The primer may be either single-stranded or double-stranded and must be sufficiently long to prime the synthesis of the desired extension product in the presence of the inducing agent. The exact length of the primer will depend upon many factors, including temperature, source of primer and use the method. For example, for diagnostic applications, depending on the complexity of the target sequence, the oligonucleotide primer typically contains 15-25 or more nucleotides, although it may contain fewer nucleotides.

The primers herein are selected to be "substantially" complementary to different strands of a particular target DNA

sequence. This means that the primers must be sufficiently complementary to hybridize with their respective strands. Therefore, the primer sequence need not reflect the exact sequence of the template. For example, a non-complementary nucleotide
5 fragment may be attached to the 5' end of the primer, with the remainder of the primer sequence being complementary to the strand. Alternatively, non-complementary bases or longer sequences can be interspersed into the primer, provided that the primer sequence has sufficient complementary with the sequence or
10 hybridize therewith and thereby form the template for the synthesis of the extension product.

As used herein, the terms "restriction endonucleases" and "restriction enzymes" refer to enzymes, each of which cut double-stranded DNA at or near a specific nucleotide sequence.

15 A cell has been "transformed" by exogenous or heterologous DNA when such DNA has been introduced inside the cell. The transforming DNA may or may not be integrated (covalently linked) into the genome of the cell. In prokaryotes, yeast, and mammalian cells for example, the transforming DNA may
20 be maintained on an episomal element such as a plasmid. With respect to eukaryotic cells, a stably transformed cell is one in which

the transforming DNA has become integrated into a chromosome so that it is inherited by daughter cells through chromosome replication. This stability is demonstrated by the ability of the eukaryotic cell to establish cell lines or clones comprised of a population of daughter
5 cells containing the transforming DNA. A "clone" is a population of cells derived from a single cell or ancestor by mitosis. A "cell line" is a clone of a primary cell that is capable of stable growth *in vitro* for many generations.

A "heterologous" region of the DNA construct is an
10 identifiable segment of DNA within a larger DNA molecule that is not found in association with the larger molecule in nature. Thus, when the heterologous region encodes a mammalian gene, the gene will usually be flanked by DNA that does not flank the mammalian genomic DNA in the genome of the source organism. In another
15 example, coding sequence is a construct where the coding sequence itself is not found in nature (e.g., a cDNA where the genomic coding sequence contains introns, or synthetic sequences having codons different than the native gene). Allelic variations or naturally-occurring mutational events do not give rise to a heterologous region
20 of DNA as defined herein.

The labels most commonly employed for these studies are radioactive elements, enzymes, chemicals which fluoresce when exposed to ultraviolet light, and others. A number of fluorescent materials are known and can be utilized as labels. These include, for example, fluorescein, rhodamine, auramine, Texas Red, AMCA blue and Lucifer Yellow. A particular detecting material is anti-rabbit antibody prepared in goats and conjugated with fluorescein through an isothiocyanate link.

Proteins can also be labeled with a radioactive element or with an enzyme. The radioactive label can be detected by any of the currently available counting procedures. The preferred isotope may be selected from ^3H , ^{14}C , ^{32}P , ^{35}S , ^{36}Cl , ^{51}Cr , ^{57}Co , ^{58}Co , ^{59}Fe , ^{90}Y , ^{125}I , ^{131}I , and ^{186}Re .

Enzyme labels are likewise useful, and can be detected by any of the presently utilized colorimetric, spectrophotometric, fluorospectrophotometric, amperometric or gasometric techniques. The enzyme is conjugated to the selected particle by reaction with bridging molecules such as carbodiimides, diisocyanates, glutaraldehyde and the like. Many enzymes which can be used in these procedures are known and can be utilized. The preferred are peroxidase, β -glucuronidase, β -D-glucosidase, β -D-galactosidase,

urease, glucose oxidase plus peroxidase and alkaline phosphatase. U.S. Patent Nos. 3,654,090, 3,850,752, and 4,016,043 are referred to by way of example for their disclosure of alternate labeling material and methods.

5 A particular assay system developed and utilized in the art is known as a receptor assay. In a receptor assay, the material to be assayed is appropriately labeled and then certain cellular test colonies are inoculated with a quantity of both the label after which binding studies are conducted to determine the extent to which the
10 labeled material binds to the cell receptors. In this way, differences in affinity between materials can be ascertained.

 An assay useful in the art is known as a "cis/trans" assay. Briefly, this assay employs two genetic constructs, one of which is typically a plasmid that continually expresses a particular receptor of
15 interest when transfected into an appropriate cell line, and the second of which is a plasmid that expresses a reporter such as luciferase, under the control of a receptor/ligand complex. Thus, for example, if it is desired to evaluate a compound as a ligand for a particular receptor, one of the plasmids would be a construct that
20 results in expression of the receptor in the chosen cell line, while the second plasmid would possess a promoter linked to the luciferase

gene in which the response element to the particular receptor is inserted. If the compound under test is an agonist for the receptor, the ligand will complex with the receptor, and the resulting complex will bind the response element and initiate transcription of the luciferase gene. The resulting chemiluminescence is then measured photometrically, and dose response curves are obtained and compared to those of known ligands. The foregoing protocol is described in detail in U.S. Patent No. 4,981,784.

The current invention is directed to a method of treating neuroectodermal tumors with a neuroectodermal tumorspecific ligand fused to a cytotoxic moiety. Possible neuroectodermal tumor targets include gliomas, meningiomas, ependymomas, medulloblastomas, neuroblastomas, gangliomas, pheochromocytomas, melanomas, PPNET's, small cell carcinoma of the lung, Ewing's sarcoma, and metastatic tumors in the brain. Preferably, the neuroectodermal tumor specific ligand is chlorotoxin fused to a cytotoxic moiety. Examples of possible cytotoxic moieties include gelonin, ricin, saponin, pseudomonas exotoxin, pokeweed antiviral protein, diphtheria toxin, and complement protein.

The current invention is also directed to a neuroectodermal tumor specific therapeutic agent in which the

neuroectodermal tumor specific ligand is an antibody against the chlorotoxin receptor believed to be a 72 kDa chloride channel. The antibody may be fused to gelonin, ricin, saponin, pseudomonas exotoxin, pokeweed antiviral protein, diphtheria toxin, or complement proteins.

The current invention is also directed to a method of differentiating neuroectodermal tumor-derived neoplastic tumor tissue from non-neoplastic tissue by incubating the tissue of interest with labeled chlorotoxin and measuring the binding of the labeled chlorotoxin, relative to normal tissue where available. The chlorotoxin may be labeled with either a fluorescent moiety or may be radiolabeled with radiolabels such as ^{131}I or ^{125}I . Fluorescent moieties can be used for detection by fluorescent microscopy or fluorescent activated cell sorting. Radiolabeled chlorotoxin can be detected by positron emission tomography scanning.

The following examples are given for the purpose of illustrating various embodiments of the invention and are not meant to limit the present invention in any fashion.

EXAMPLE 1

Summary of Chlorotoxin Results from Glioma Experiments

Recent studies demonstrated a common antigen is
5 expressed by the vast majority of glioma cells. This antigen is
targeted by chlorotoxin (Ctx or TM-601), a 36 amino acid peptide
originally isolated from *Leiurus quinquestriatus* scorpion venom.
Chlorotoxin selectively binds to the membrane of glioma cells
allowing selective targeting of these cells within the CNS (18). The
10 antigen targeted by this peptide appears to be a chloride ion channel
although the antigen has not yet been unequivocally identified at the
molecular level. Thus far, the data indicates that chlorotoxin binds to
a membrane protein of 72 kDa molecular weight that is
preferentially expressed in the cytoplasmic membrane of glioma cell.
15 Binding of the peptide enhances glioma cell proliferation (19) and
inhibits the ability of glioma cells to migrate in Transwell assays, an
in vitro assay to evaluate tumor invasiveness (20). Chlorotoxin
appears to exert these effects by reducing the membrane
permeability to Cl⁻ ions thereby preventing cell volume changes that
20 are required to allow cells to invade healthy tissue (20). Thus, the

most likely action of chlorotoxin is on a glioma chloride channel previously extensively characterized (19).

5

EXAMPLE 2

Immunohistochemical Staining of Gliomas with Chlorotoxin

Over 250 frozen or paraffin sections of human biopsy tissues were histochemically stained with a chemically synthesized form of chlorotoxin containing a detectable biotin group chemically attached to the N terminus (TM-601). Binding of the TM-601molecule was observed on selective cells associated with the essentially all glioma tumors with up to 95% positive cells per tumor. Based on these studies, it has been proposed to utilize chlorotoxin as a glioma specific marker and as a potential therapeutic tool for targeting glioma tumors. For such purposes, chlorotoxin linkage of radioactive molecules or cytotoxic moieties such as saporin could be employed.

20

EXAMPLE 3

Recombinant DNA Manipulation of Chlorotoxin

Using techniques well known in the art, one may prepare
5 recombinant proteins specifically engineered to mimic the binding
and action of the native toxin. The biological activity of the synthetic
chlorotoxin is as effective for chloride ion channel blockade as the
native venom toxin. Recombinant techniques are used to synthesize
chlorotoxin in *E coli* using a modified PGEX vector system and the
10 toxin may be linked to various fusion proteins using common
restriction sites. After synthesis of recombinant chlorotoxin, it may
be linked to various cytotoxic fusion proteins including glutathione-S-
transferase (GST), gelonin, ricin, diphtheria toxin, complement proteins
and radioligands and other such proteins as are well known in the
15 immunotoxin art.

EXAMPLE 4

Antibodies against the chlorotoxin-binding chloride ion channel

20 Antibodies to the chloride ion channels in glial-derived
tumors may be prepared as follows. Polyclonal antisera are

generated by injecting fusion proteins created between the glutathione-S-transferase and the chlorotoxin insert into mice or rabbits. Mice are immunized with 0.5 ml of a 1:1 emulsion of 1 mg/ml purified fusion protein in Freund's complete adjuvant and subsequently with two additional injections after 14 and 28 days in Freund's incomplete adjuvant. The mouse and rabbit antibodies are purified from the antisera using the GST fusion protein immobilized on nitrocellulose filters. The antibodies are then examined for binding specificity in various tissues.

EXAMPLE 5

Rationale for the examination of neuroectodermally derived tumors for chlorotoxin binding

Given the similarities that can be shared between gliomas and other neuroectodermally-derived cells, and the arguments developed that propose similar propensities of neuroectodermal cells to migrate, a thorough investigation was undertaken to examine neuroectodermally derived tumors for the expression of chlorotoxin binding sites.

EXAMPLE 6

Preparation of sections from frozen or paraffin-embedded human 5 biopsies

Most of the samples of human tissue, from both sexes, all ages and race were obtained through the Cooperative Human Tissue Network, Tissue Procurement at UAB, UAB hospitals and the Human Brain Tissue Bank in London, Canada. Snap frozen tissue and fresh
10 tissue embedded in a freezing gel were sliced at 8 microns and picked up onto positively charged glass slides. The sections were then fixed in 4% paraformaldehyde or millonig according to the staining protocol. Paraffin blocks were sectioned and prepared according to standard procedures.

EXAMPLE 7

Examination of biopsy samples for chlorotoxin binding

20 Biopsy sections were blocked for 1 hour in 10% normal goat serum in PBS and treated with a dilution of biotinylated

chlorotoxin overnight at 4°C. After thorough rinsings, the stainings were developed by avidin-biotin complex (ABC) technique (Vectastain Elite ABC Kit from Vector Laboratories, Burlington, CA) and visualized by the colorimetric reaction of DAB (3'3'-diaminobenzidine; Vector Laboratories) with the ABC complex.

The biopsy sections were counterstained with methyl green, a nuclear dye, to more easily visualize the unstained cells. Non-specific background label can vary from experiment to experiment due to changes in the effective concentrations of the label, condition of the tissue or the duration of the reaction. Therefore, a control section was identically stained with methyl green but without the biotinylated chlorotoxin. Positive cell staining is identified by chlorotoxin-labeling above background when compared to its individual control of a successive slice. Cells containing high amounts of endogenous peroxidase exhibit dark background staining in the controls due to the reaction of DAB with the peroxidases.

Finally, a third adjacent section was stained with both hematoxylin, a cell nuclei specific stain, and eosin, a cytoplasmic stain. Therefore, for each tissue analyzed, three adjacent section were stained. These are shown in photomicrographs providing

evidence of the specificity of TM-601 chlorotoxin binding to tumors of neuroectodermal derivation in comparison to controls. In the photomicrographs, adjacent sections are identified as follows: TM-601: biotinylated chlorotoxin detected by a brown reaction product of DAB with the biotin and further counterstained with methyl green; Control: the control section stained with only methyl green; and, H&E: the hematoxylin and eosin stained section.

EXAMPLE 8

Glioblastoma multiforme (GBM) tumors

Glioblastoma multiforme (GBM) were stained with the biotinylated chlorotoxin (TM-601). These tumors are extremely reactive to biotinylated chlorotoxin as 25 out of 25 patient samples tested positive as seen in Figure 1. This glioblastoma multiforme can be compared to the staining of the normal human brain tissue with biotinylated chlorotoxin (18/23 negative). Figure 2 shows a representative staining. Normal brain tissue demonstrates a lack of TM-601 staining. This is consistent with earlier evidence of specificity of chlorotoxin binding to gliomas.

EXAMPLE 9

GFAP staining in the normal brain tissue

5 The biopsy section was blocked for 1 hour in 10% normal
goat serum in PBS and then stained with antibodies against glial
fibrillary acidic protein (GFAP; DAKO corporation, Carpinteria CA)
overnight. The secondary antibody conjugated to peroxidase was
applied to the rinsed tissue for 2 hours, rinsed again before the stain
10 was visualized with DAB. A typical glial fibrillary acidic protein stain
of normal brain is shown in Figure 2. Normal brain was positive for
glial fibrillary acidic protein staining where it stained the astrocytes
typically present in normal tissue.

EXAMPLE 10

Neuroblastomas

Neuroblastomas are a tumor primarily found in children
20 with a high incidence in the adrenals. Neuroblastoma show TM-601

reactivity above the control staining as seen in Figure 3. Six out of seven neuroblastomas were positive for chlorotoxin binding.

5

EXAMPLE 11

Pheochromocytomas

Pheochromocytomas are neoplastic chromaffin cells of the adrenal glands. This tumor also show a high degree of staining as seen in Figure 4. Five out of six pheochromocytomas were positive for staining with biotinylated chlorotoxin, especially in comparison to TM-601 staining of the normal adrenals (3/3 negative) seen in Figure 5.

EXAMPLE 12

Melanomas

Figure 6 shows the biotinylated chlorotoxin staining of a melanoma metastasized to the brain. Seven out of seven melanoma brain metastasis were positive for TM-601. In addition, melanoma

metastasized to the lung were analyzed as seen in Figure 7. Normal skin, however, is unreactive to TM-601 (6/6 negative) (Figure 8) although there is some background staining in the melanocytes even in the controls.

5

EXAMPLE 13

Small cell lung carcinomas

10 Small cell lung carcinomas are reactive to TM-601. There is good contrast between the cells that stain and those that do not (Figure 9). The cells positive for TM-601 in the control (middle panel) are red blood cells which present high levels of background peroxidase stain. This TM-601 specificity can be further
15 demonstrated by comparing the TM-601 staining of the small cell carcinoma (2/3 positive) and the normal lung (3/3 negative) (Figure 10).

20

EXAMPLE 14

Medulloblastomas

Another neuroectodermally derived tumor type are the
5 medulloblastomas. They exhibit specific reactivity to TM-601 as
seen in Figure 11 (4/4 positive).

EXAMPLE 15

Ewing's sarcoma

Ewing's sarcoma, a rare bone cancer sometimes found in
soft tissue, is TM-601 positive (2/2) (Figure 12).

EXAMPLE 16

Testing of potential sites of chlorotoxin administration for side effects

To aid in the design of drug therapy with this product,
20 various normal tissues were stained with TM-601 to determine
possible sites where side effects may occur. Preliminary evidence

indicates that some of the most common targets for side effects such as the stomach and liver, are TM-601 negative (2/2 negative samples for both tissues, thus far) (Figures 13 and 14 respectively).

The staining of spleen tissue is also shown in Figure 15 (3/3

5 negative). The TM-601 staining of other normal human tissues is summarized in Table 1.

Table I

Tumor or Tissue Type	# of Cases	Chlorotoxin-binding
Primary Brain Tumors:		
Gliomas:		
WHO grade IV: glioblastoma multiforme	25	Positive
WHO grade III: anaplastic astrocytoma	2	Positive
WHO grade II: low grade	2	Positive
WHO grade I: pilocytic astrocytoma	11	Positive
Oligodendrogliomas	6	Positive
Other gliomas	3	Positive
Gangliomas	3	Positive
Meningiomas	18	Positive
Ependymomas	3	Positive
Other Primary Brain Tumors:		
Epidermoid cysts in brain	3	2/3 Positive
Brain tumors-unknown pathology	15	14/15 Positive
Pituitary gland of GBM patient	2	Positive
Secondary Brain Tumors:		
Metastatic tumors to brain	14	12/14 Positive
Comparison of Brain Tissues		
Alzheimer brains	8	Negative
Brain, normal or uninvolved	24	18/24 Negative
Epilepsy/gliosis/stroke	6	Positive
Uninvolved brain of GBM	3	Negative (autopsy)
Tumors of Neuro Ectodermal Origin:		
Medulloblastoma	4	Positive
Neuroblastoma	7	6/7 Positive
Ganglioneuroma	4	Positive
Melanomas	7	Positive
Pheochromocytoma	6	5/6 Positive
PPNET	1	Negative
Small cell carcinoma lung	3	2/3 Positive
Ewing's sarcoma	2	Positive

Table I (Continued)

Tumor or Tissue Type	# of Cases	Chlorotoxin-binding
Other Human Tissues		
Colon	3	Negative
Endometrium/myometrium	2	Negative
Heart	2	Negative
Kidney	2	Cortex is Positive/ Medulla is Negative
Adrenal Gland	3	Negative
Liver	3	Negative
Lung	3	Negative
Lung, small cell carcinoma	1	Positive
Meninges	3	Negative
Muscle, skeletal	2	Negative
Pancreas, fibrosis	1	Negative
Ovary	2	Mostly Negative/ a few cells are positive
Skin, from thigh, abdomen or breast	6	Negative
Spleen	3	Negative
Stomach	2	Negative
Testes	2	Mostly negative/ a few cells are positive
Thyroid	1	Negative

EXAMPLE 17

Summary of Tested Tumors and Tissues

As summarized in Table 1, the vast majority of neuroectodermally derived tumors bind chlorotoxin, indicating that chlorotoxin has a more widespread utility to target tumors of neuroectodermal origin. Specifically, primitive neuroectodermal tumor tumors have been tested from 34 patients, 31 of which showed chlorotoxin specificity in the tumor material as seen in Table 1. This staining was compared with the chlorotoxin staining of other types of CNS and PNS tumors as well as the comparison to various normal human tissues.

TM-601 specifically associates with neuroectodermally-derived tumors including medulloblastomas, neuroblastomas, ganglioneuromas, melanomas, pheochromocytomas, small cell lung carcinomas and Ewing's sarcomas. Thus, chlorotoxin-derived molecules can be utilized to target specifically for therapeutic or diagnostic purposes the above identified neuroectodermally-derived tumors. Likewise, these tumors can also be targeted by other molecules such as antibodies that bind to the chlorotoxin receptor, presumed to be the 72kD Cl^- ion channel.

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Any patents or publications mentioned in this specification are indicative of the levels of those skilled in the art to

which the invention pertains. These patents and publications are herein incorporated by reference to the same extent as if each individual publication was specifically and individually indicated to be incorporated by reference.

5 One skilled in the art will readily appreciate that the present invention is well adapted to carry out the objects and obtain the ends and advantages mentioned, as well as those inherent therein. The present examples along with the methods, procedures, treatments, molecules, and specific compounds described herein are
10 presently representative of preferred embodiments, are exemplary, and are not intended as limitations on the scope of the invention. Changes therein and other uses will occur to those skilled in the art which are encompassed within the spirit of the invention as defined by the scope of the claims.

WHAT IS CLAIMED IS:

1. A method of treating an individual having a neuroectodermal tumor, comprising the step of:

5 administering a pharmaceutical composition comprising a pharmaceutically effective dose of a neuroectodermal tumor specific ligand fused to a cytotoxic moiety and a pharmaceutically acceptable carrier.

10 2. The method of claim 1, wherein the neuroectodermal tumor is a tumor type treated is selected from the group consisting of ependymomas, medulloblastomas, neuroblastomas, gangliomas, pheochromocytomas, melanomas, peripheral primitive
15 neuroectodermal tumors, small cell carcinoma of the lung, Ewing's sarcoma, and metastatic tumors in the brain.

20 3. The method of claim 1, wherein the neuroectodermal tumor specific ligand is chlorotoxin.

4. The method of claim 3, wherein said chlorotoxin is selected from the group consisting of native chlorotoxin, synthetic chlorotoxin and recombinant chlorotoxin.

5

5. The method of claim 1, wherein said cytotoxic moieties is selected from the group consisting of gelonin, ricin, saponin, pseudomonas exotoxin, pokeweed antiviral protein, diphtheria toxin, and complement proteins.

10

6. The method of claim 1, wherein the neuroectodermal tumor specific ligand is an antibody against the chlorotoxin receptor.

15

7. A method of differentiating neuroectodermal tumor-derived neoplastic tumor tissue from non-neoplastic tissue, comprising the steps of:

contacting a tissue of interest with labeled chlorotoxin
which binds specifically to neuroectodermal tumor neoplastic tumor
tissue; and

measuring the binding of the labeled chlorotoxin, wherein
5 an elevated level of binding, relative to normal tissue, indicates that
the tissue is neoplastic.

8. The method of claim 7, wherein said chlorotoxin is
10 labeled with a detection moiety.

9. The method of claim 8, wherein said detection
moiety is selected from the group consisting of a fluroschrome, biotin,
15 a colorimetric agent linked to an enzyme substrate.

10. The method of claim 8, wherein said labeled
chlorotoxin binding is determined by a method selected from the
20 group consisting of fluorescent microscopy, ELIZA and fluorescent
activated cell sorting.

11. The method of claim 7, wherein said labeled chlorotoxin is radiolabeled.

5

12. The method of claim 11, wherein said radiolabeled chlorotoxin is selected from the group consisting of ^{131}I -chlorotoxin and ^{125}I -chlorotoxin.

10

13. The method of claim 7, wherein the level of radiolabeled chlorotoxin binding affinity indicative of neoplastic tissue is from about 5 nanomolar to about 5 micromolar.

15

14. The method of claim 13, wherein said labeled chlorotoxin binding is determined using positron emission tomography scanning.

ABSTRACT OF THE DISCLOSURE

The present invention provides fusion proteins for the
5 detection and treatment of neuroectodermal tumors. Previous work
demonstrated that chlorotoxin is specific for glial-derived or
meningioma-derived tumor cells. The current invention has
extended the use of chlorotoxin-cytotoxin fusion proteins to treat the
whole class neuroectodermal tumors such as gliomas, meningiomas,
10 ependymomas, medulloblastomas, neuroblastomas, gangliomas,
pheochromocytomas, melanomas, PPNET's, small cell carcinoma of the
lung, Ewing's sarcoma, and metastatic tumors in the brain. Also,
diagnostic methods are provided for screening neoplastic
neuroectodermal tumors.

Normal Brain

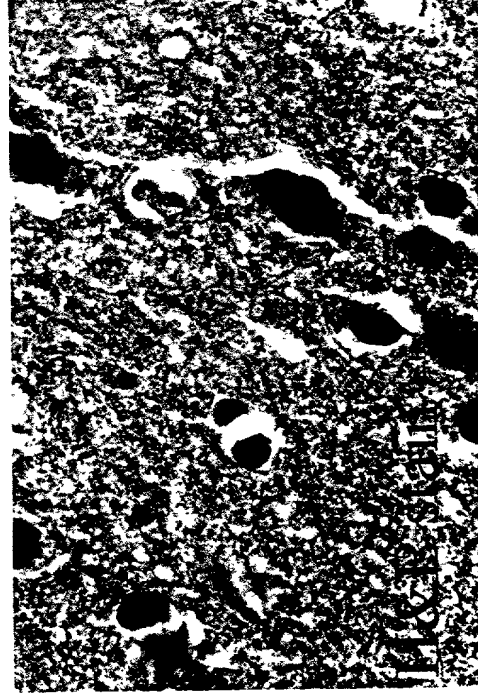
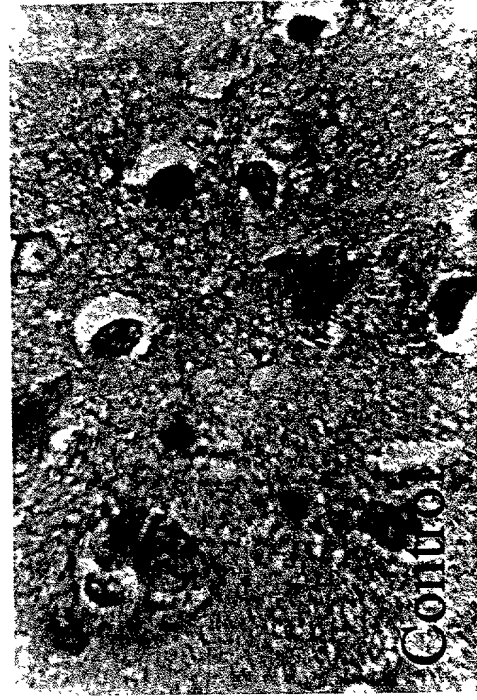
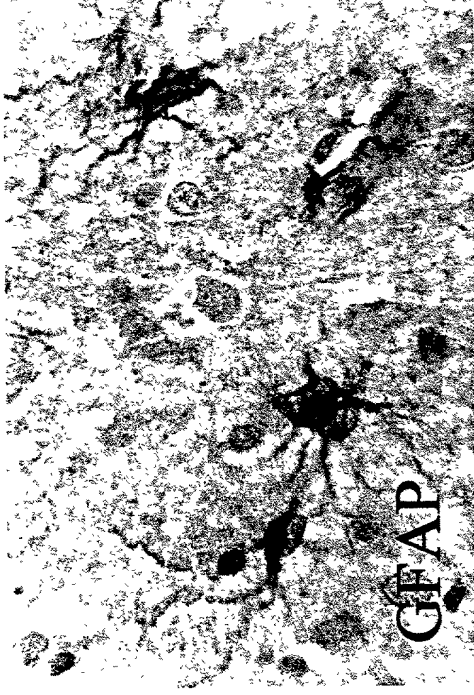
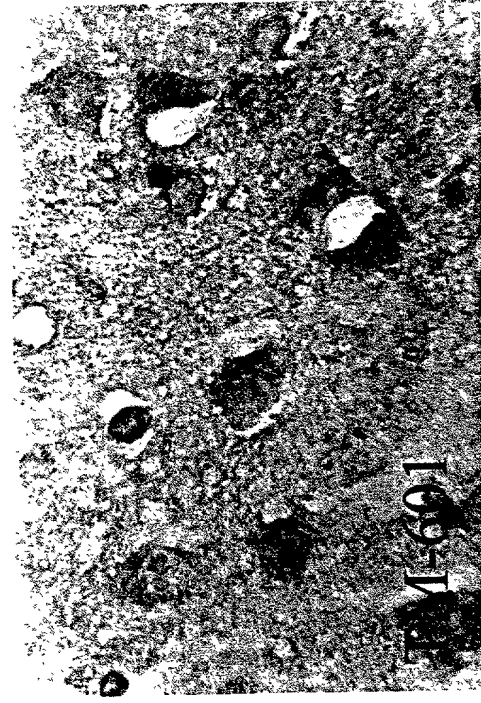
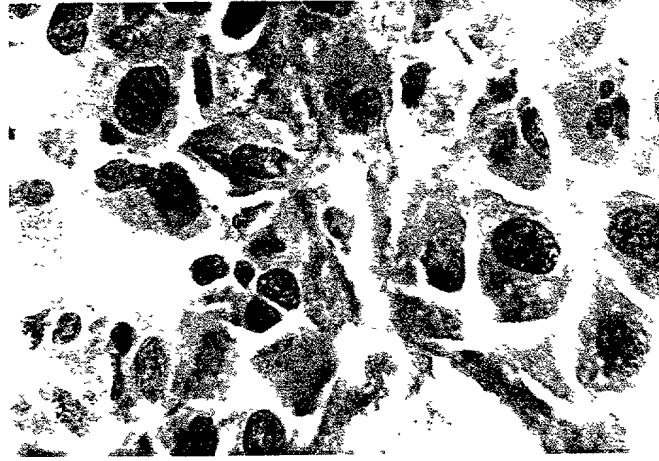


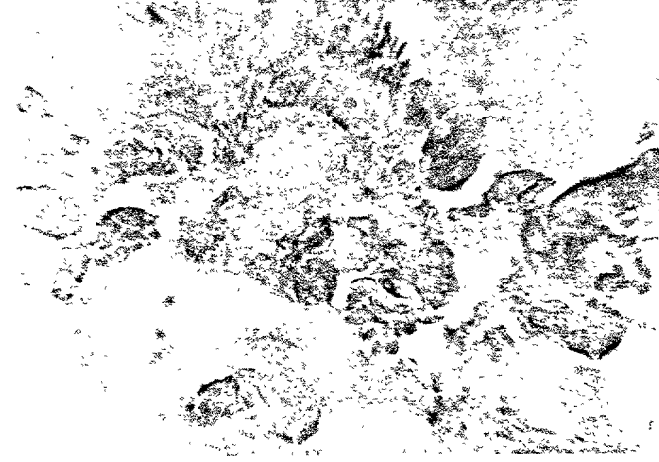
FIGURE 1

TM-601 Specific Binding to PNET tumors
S.A. Lyons and H.W. Sontheimer

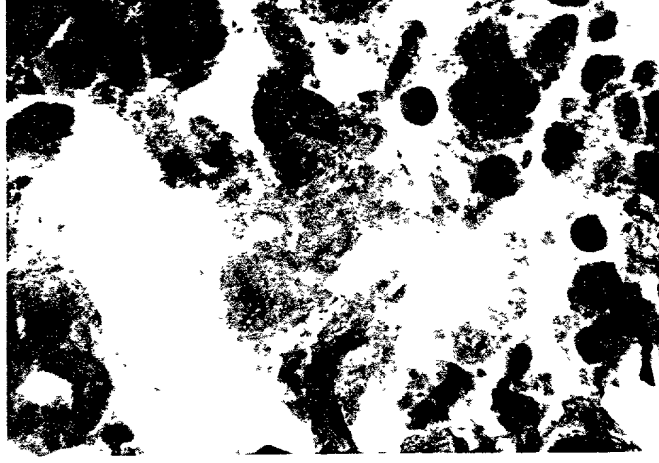
Glioblastoma multiforme



TM-601



Control

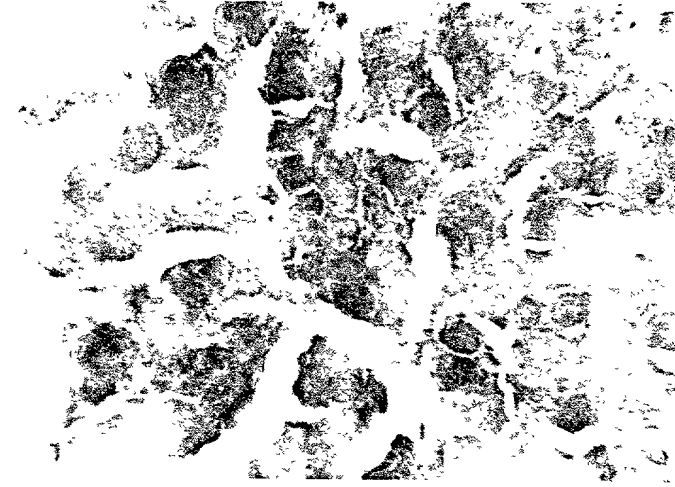


H&E stain

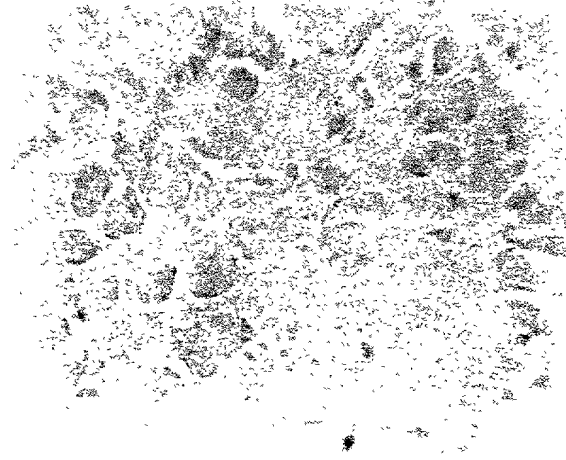
FIGURE 2

TM-601 Specific Binding to PNET tumors
S.A. Lyons and H.W. Sontheimer

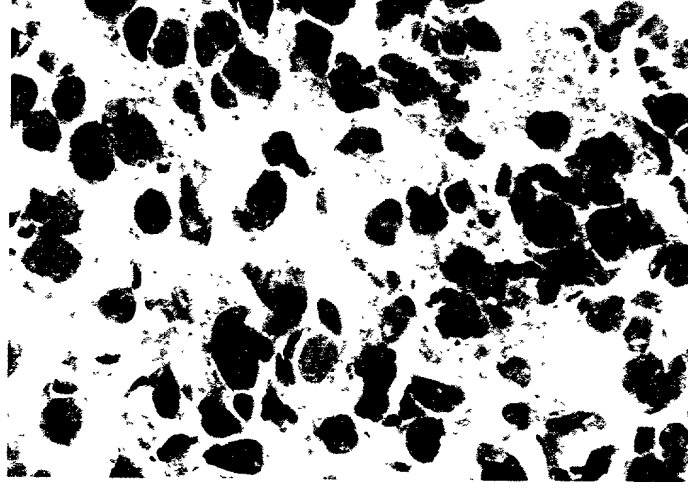
Adrenal Mass Neuroblastoma



TM-601



Control

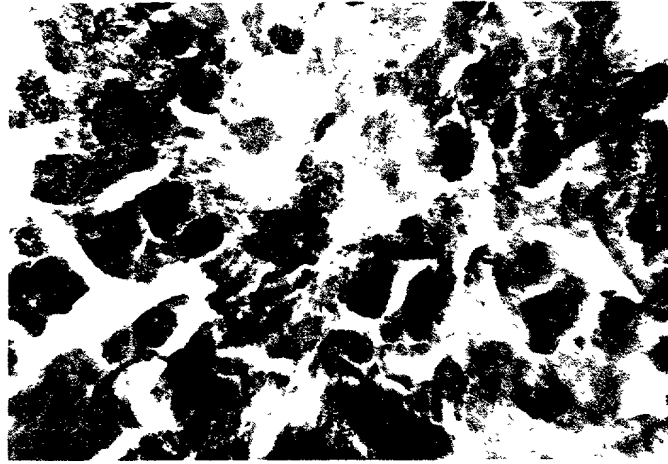


H&E stain

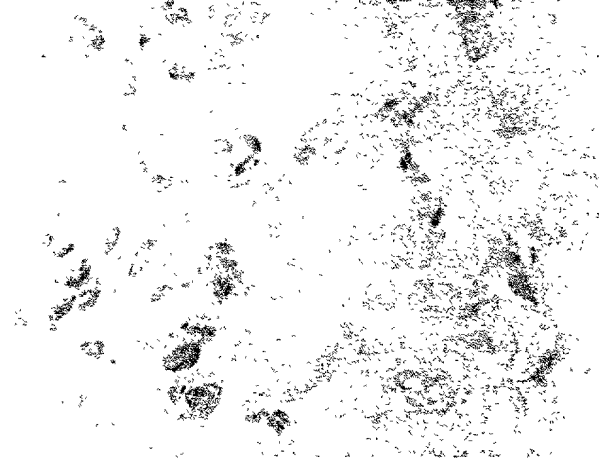
FIGURE 3

TM-601 Specific Binding to PNET tumors
S.A. Lyons and H.W. Sonthmeier

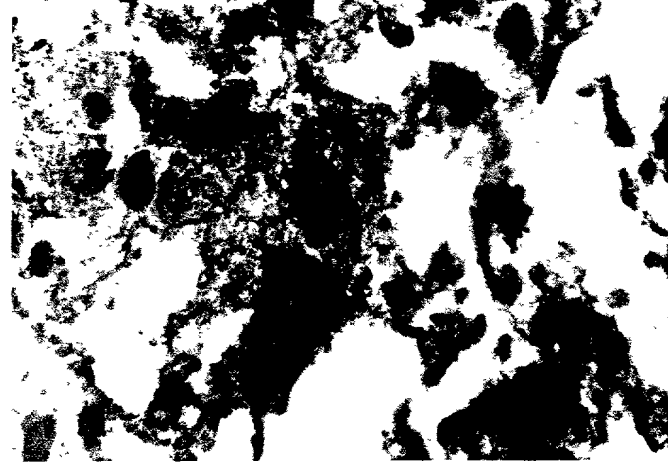
Pheochromocytoma



TM-601



Control

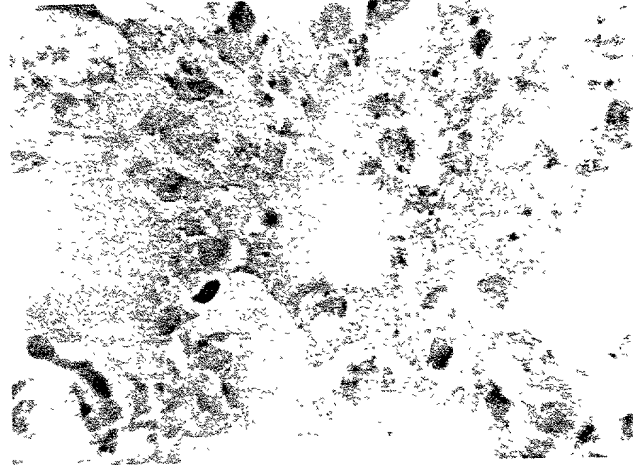


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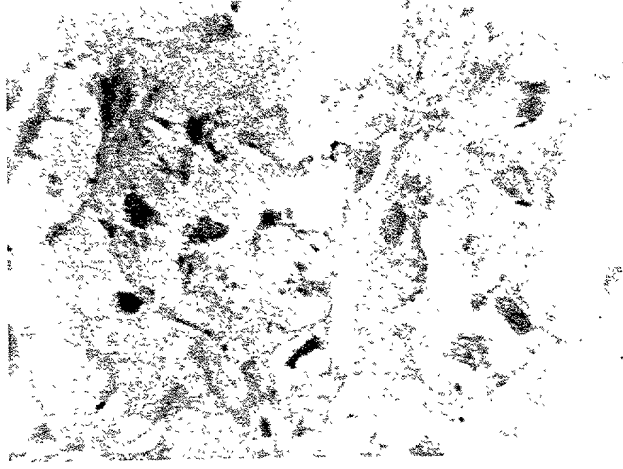
FIGURE 4

TM-601 Specific Binding to PNET tumors
S.A. Lyons and H.W. Sontheimer

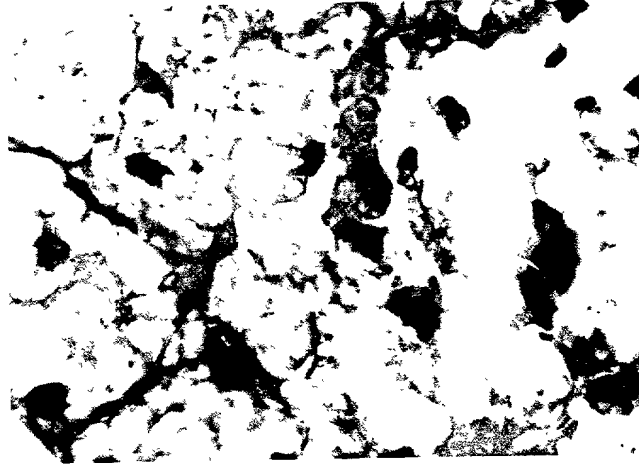
Normal Adrenal



TM-601



Control

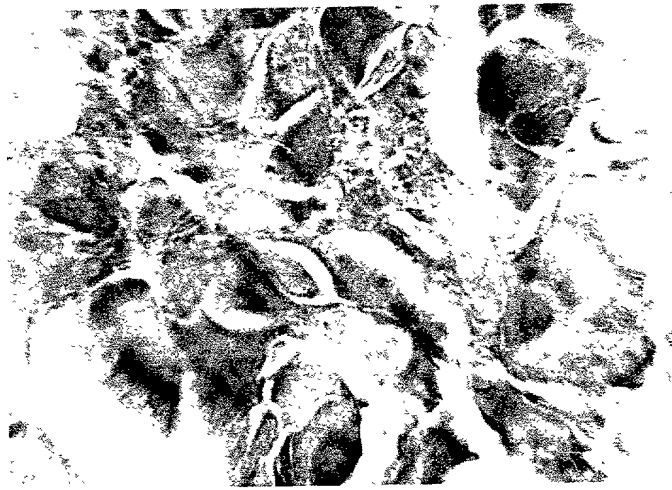


H&E stain

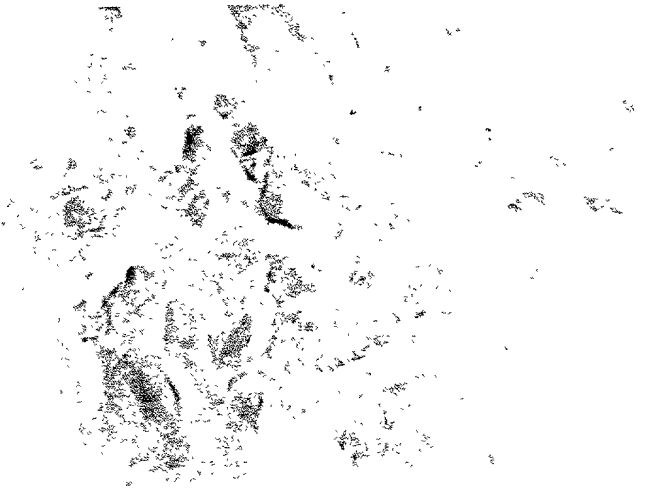
FIGURE 5

TM-601 Specific Binding to PNH Tumors
S.A. Lyons and H.W. Sontheimer

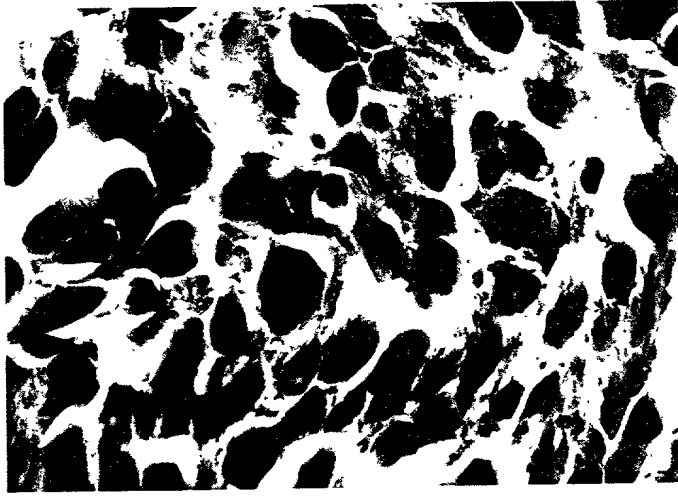
Melanoma to Brain



TM-601



Control

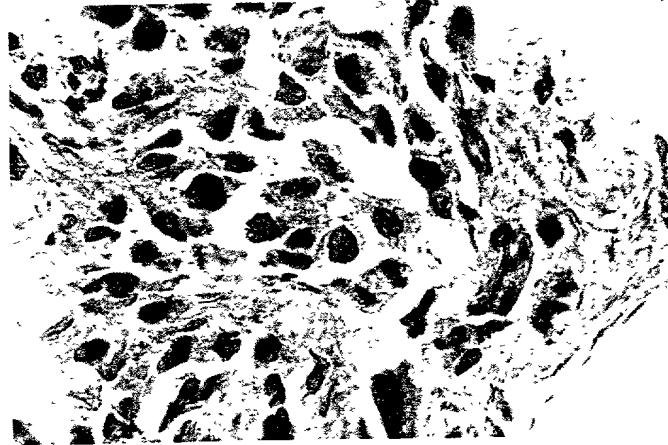


H&E stain

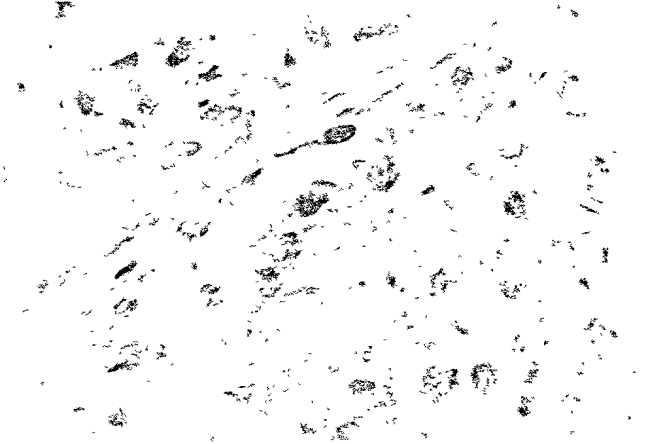
FIGURE 6

TM-601 Specific Binding to PNL1 tumors
S.A. Lyons and H.W. Sonthelmer

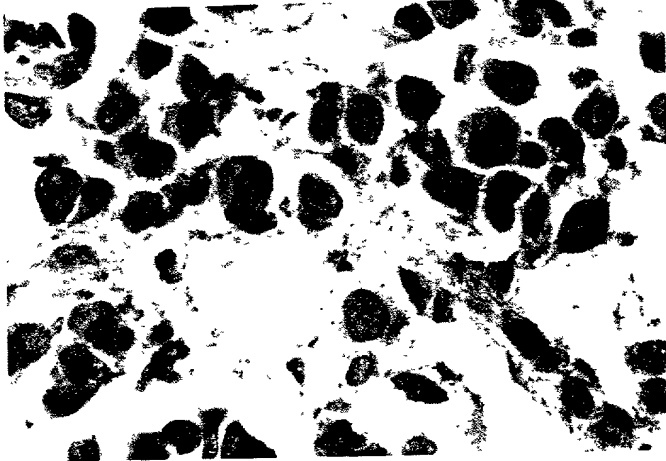
Melanoma to Lung



TM-601



Control



H&E stain

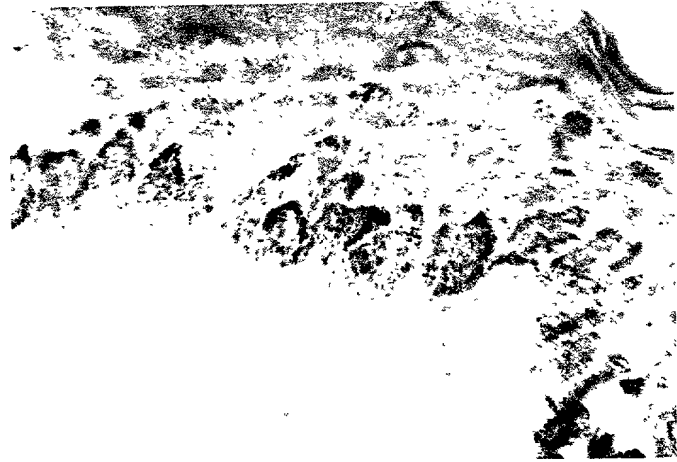
FIGURE 7

TM-601 Specific Binding to PNL-I tumors
S.A. Lyons and H.W. Sontheimer

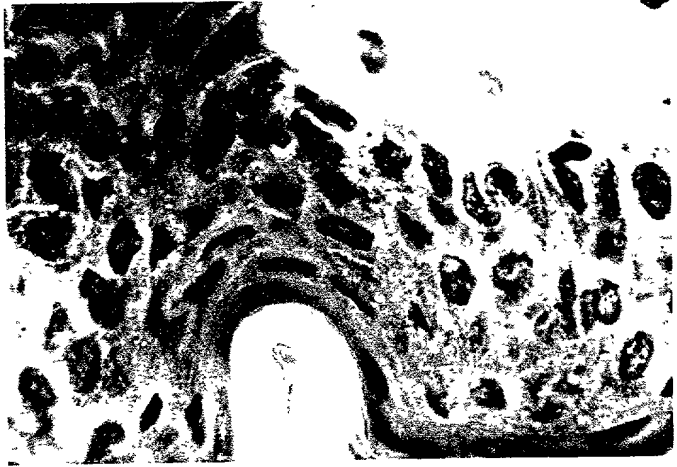
Normal Breast Skin



TM-601



Control

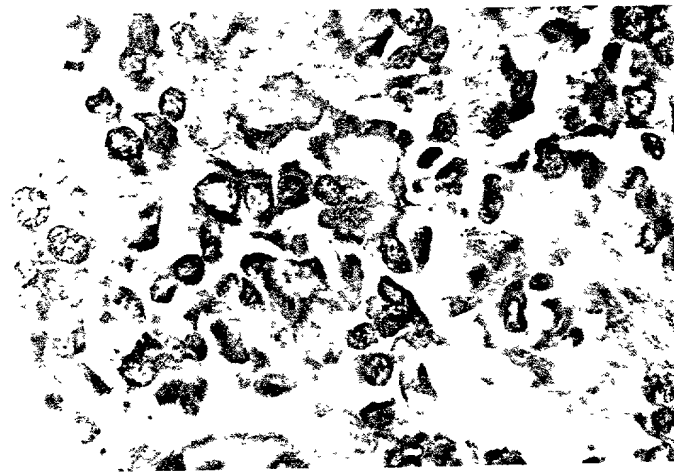


H&E stain

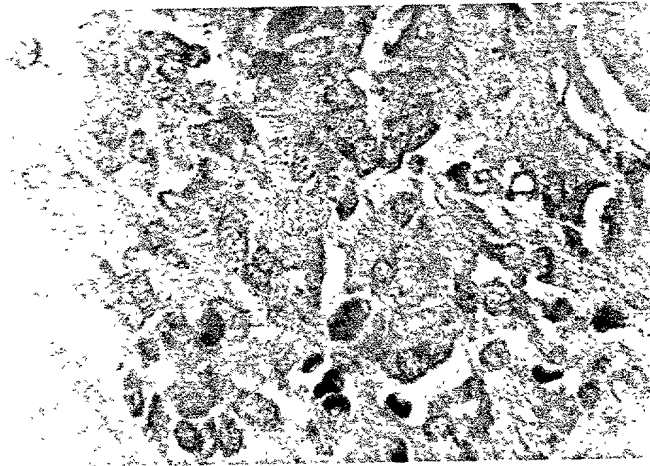
FIGURE 8

TM-601 Specific Binding to PNET tumors
S.A. Lyons and H.W. Sontheimer

Small Cell Lung Carcinoma



TM-601



Control



H&E stain

FIGURE 9

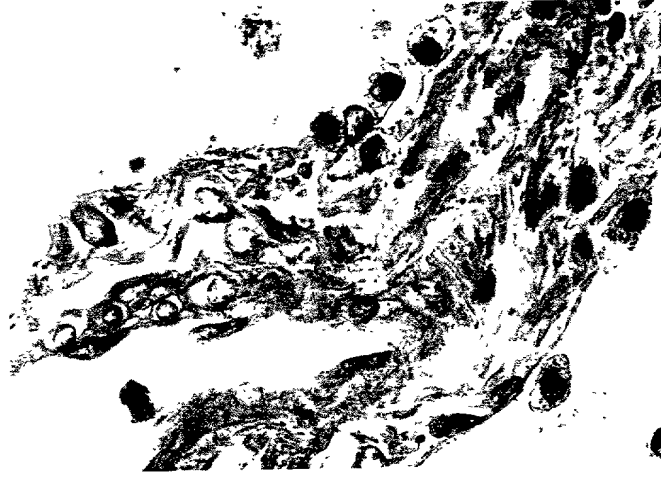
TM-601 Specific Binding to PNET tumors
S.A. Lyons and H.W. Sontheimer

Normal Lung



TM-601

Control

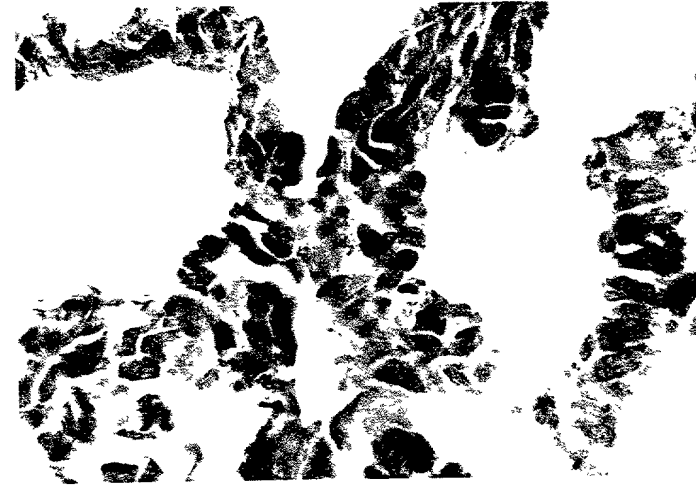


H&E stain

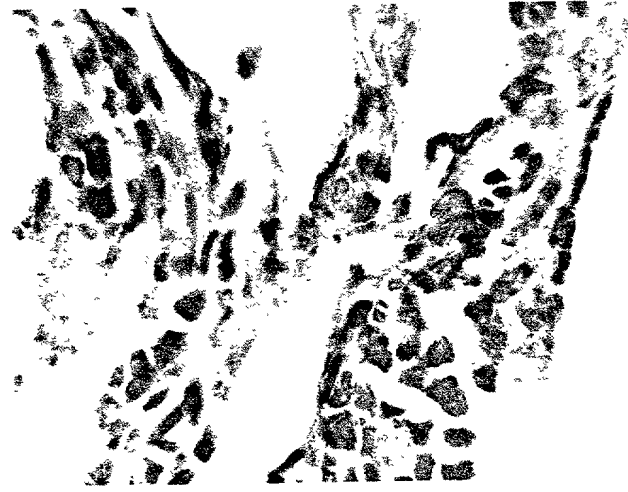
FIGURE 10

TM-601 Specific Binding to PNL-1 tumors
S.A. Lyons and H.W. Sontheimer

Medulloblastoma



TM-601



Control

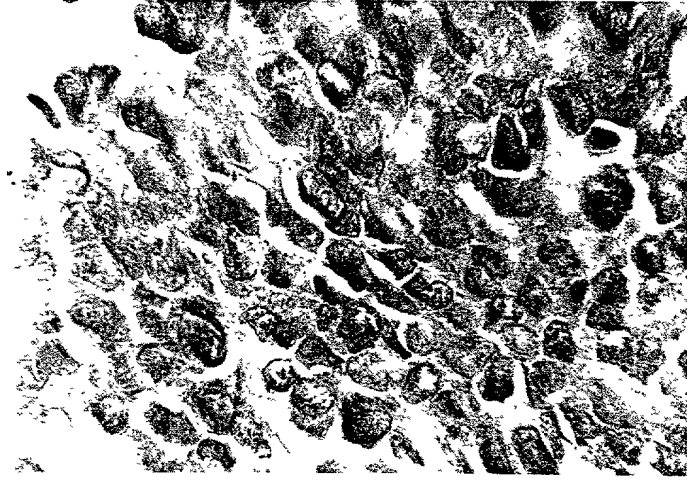


H&E stain

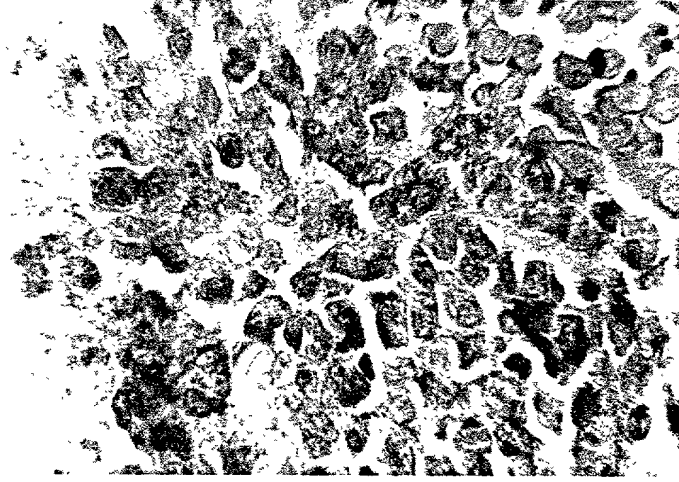
FIGURE 11

TM-601 Specific Binding to PNET tumors
S A Lyons and H W. Sontheimer

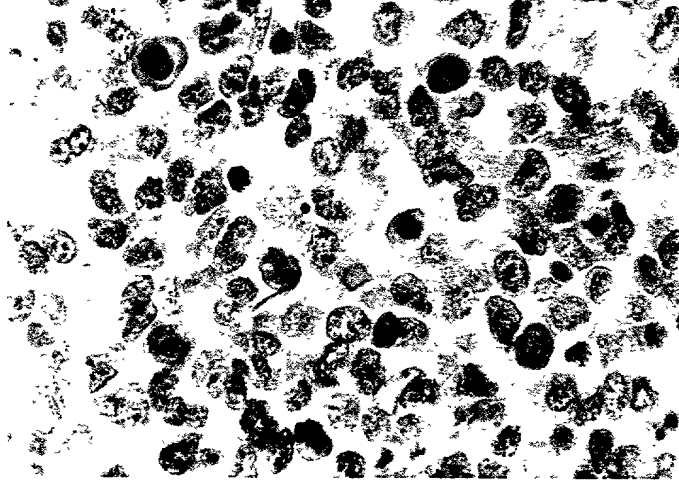
Ewing's Sarcoma



TM-601



Control

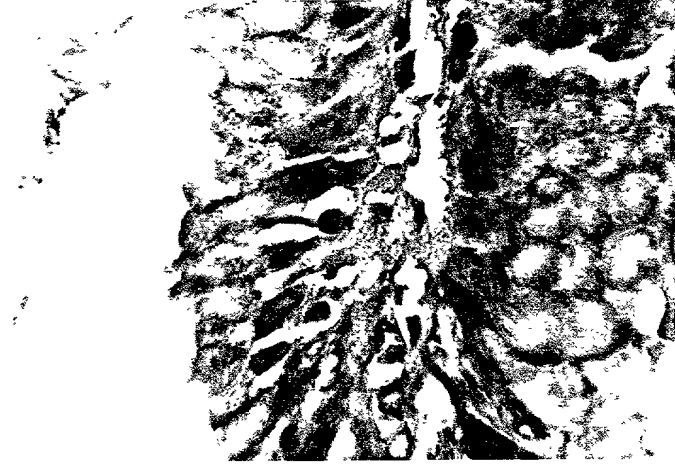


H&E stain

FIGURE 12

TM-601 Specific Binding to PNF T tumors
S.A. Lyons and H.W. Sontheimer

Normal Stomach



TM-601



Control



H&E stain

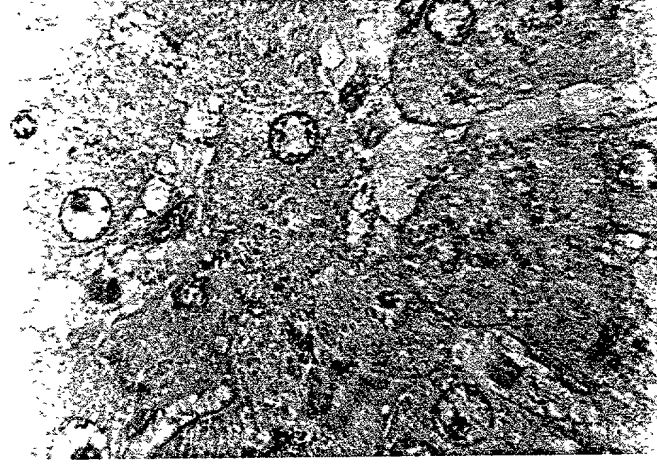
FIGURE 13

TM-601 Specific Binding to PNI, I tumors
S.A. Lyons and H.W. Sontheimer

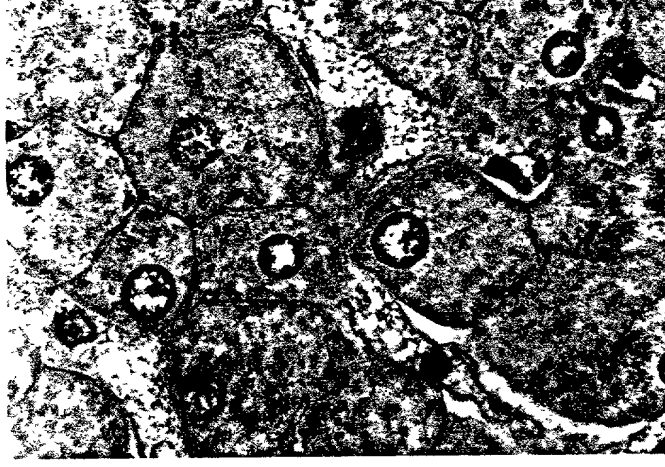
Normal Liver



TM-601



Control

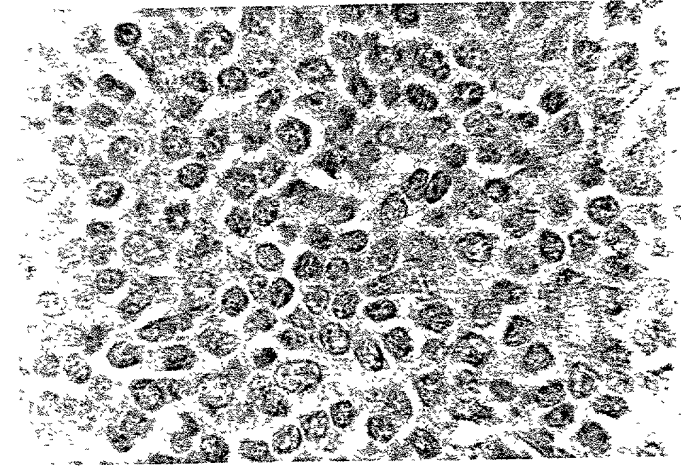


H&E stain

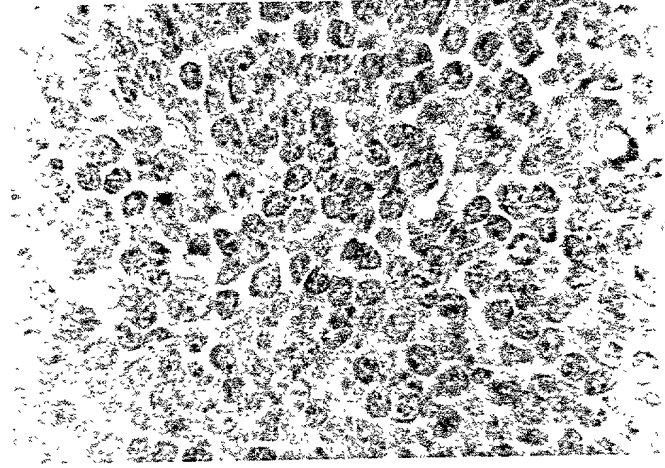
FIGURE 14

TM-601 Specific Binding to PNL1 tumors
S A Lyons and H W Sontheimer

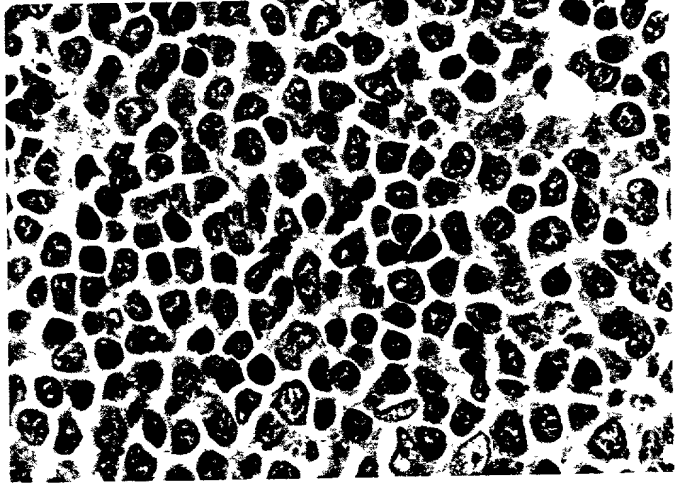
Normal Spleen



TM-601



Control



H&E stain

FIGURE 15

TM-601 Specific Binding to PNI T tumors
S.A. Lyons and H.W. Sontheimer

FROM : MCGREGOR&ADLER, P.C.

PHONE NO. :

Apr. 21 1999 11:51AM P2

DOCKET NO: D6218

COMBINED DECLARATION AND POWER OF ATTORNEY

I, **Susan A. Lyons**, hereby declare that:

My residence, post office address and citizenship are as stated below next to my name; I believe I am the original, first and co-inventor, together with **Harald W. Sontheimer**, of the subject matter which is claimed and for which a patent is sought on the invention entitled, **Diagnosis and Treatment of Neuroectodermal Tumors**; the specification of which is attached hereto.

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above. I acknowledge the duty to disclose all information I know to be material to patentability in accordance with Title 37, Code of Federal Regulations, 37 CFR §1.56(a).

I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith: **Dr. Benjamin Adler**, Registration No. 35,423. Address all telephone calls to **Dr. Benjamin Adler** at telephone number 713/777-2321. Address correspondence to **Dr. Benjamin Adler**, MCGREGOR & ADLER, L.L.P., 8011 Candle Lane, Houston, TX 77071.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patents issued thereon.

Full Name of Inventor: Susan A. Lyons Ph.D.

Inventor's Signature:  Date: 4.21.99

Residence Address: 2848 Highland Ave. S. #3; Birmingham, AL 35205

Citizen of: The United States of America

Post Office Address: Birmingham, AL 35205

667240-TE096260

FROM : MCGREGOR&ADLER, P.C.

PHONE NO. :

Apr. 21 1999 11:58AM P2

DOCKET NO: D6218

COMBINED DECLARATION AND POWER OF ATTORNEY

I, **Harald W. Sontheimer**, hereby declare that:

My residence, post office address and citizenship are as stated below next to my name; I believe I am the original, first and co-inventor, together with **Susan A. Lyons**, of the subject matter which is claimed and for which a patent is sought on the invention entitled, **Diagnosis and Treatment of Neuroectodermal Tumors**; the specification of which is attached hereto.

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above. I acknowledge the duty to disclose all information I know to be material to patentability in accordance with Title 37, Code of Federal Regulations, 37 CFR §1.56(a).

I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith: **Dr. Benjamin Adler**, Registration No. 35,423. Address all telephone calls to **Dr. Benjamin Adler** at telephone number 713/777-2321. Address correspondence to **Dr. Benjamin Adler**, MCGREGOR & ADLER, L.L.P., 8011 Candle Lane, Houston, TX 77071.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patents issued thereon.

Full Name of Inventor: Harald W. Sontheimer

Inventor's Signature: [Signature] Date: 4/21/99

Residence Address: 1704 Russet Woods Lane. Birmingham, AL 35294

Citizen of: Germany

Post Office Address: Birmingham, AL 35294

661240-TE096260